

**Amendments to the Claims:**

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This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Original) A pharmaceutical composition comprising a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous protein or/and a functional fragment thereof, a nucleic acid molecule encoding a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous protein or/and a functional fragment thereof or/and a modulator/effector of said nucleic acid molecule or/and said protein, preferably together with pharmaceutically acceptable carriers, diluents or/and additives.
2. (Original) The composition of claim 1, wherein the nucleic acid molecule is a vertebrate or insect *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* nucleic acid, particularly encoding a human protein as described in Table 1, or/and a nucleic molecule which is complementary thereto or/and a functional fragment thereof or/and a variant thereof.
3. (Currently Amended) The composition of claim 1 or 2, wherein said nucleic acid molecule is selected from the group consisting of
  - (a) a nucleic acid molecule encoding a polypeptide as shown in Table 1 or/and an isoform, fragment, or/and variant of said polypeptide;

- (b) a nucleic acid molecule which comprises or is the nucleic acid molecule as shown in Table 1;
- (c) a nucleic acid molecule being degenerate with as a result of the genetic code to the nucleic acid sequences as defined in (a) or (b),
- (d) a nucleic acid molecule that hybridizes at 50°C in a solution containing 1 x SSC and 0.1% SDS to a said nucleic acid molecule ~~as defined in claim 2~~ or/and as defined in (a) to (c) or/and a nucleic acid molecule which is complementary thereto;
- (e) a nucleic acid molecule that encodes a polypeptide which is at least 85%, preferably at least 90%, more preferably at least 95%, more preferably at least 98% and up to 99,6% identical to the human *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous protein, preferably as described in Table 1 ~~or as defined in claim 2~~ or to a polypeptide as defined in (a);
- (f) a nucleic acid molecule that differs from the nucleic acid molecule of (a) to (e) by mutation and wherein said mutation causes an alteration, deletion, duplication or premature stop in the encoded polypeptide.

4. (Currently Amended) The composition of ~~any one of claims 1-3~~ claim 1, wherein the nucleic acid molecule is a DNA molecule, particularly a cDNA or a genomic DNA.

5. (Currently Amended) The composition of ~~any one of claims 1-4~~ claim 1, wherein said nucleic acid encodes a polypeptide contributing to regulating the energy homeostasis or/and the metabolism of triglycerides.
6. (Currently Amended) The composition of ~~any one of claims 1-5~~ claim 1, wherein said nucleic acid molecule is a recombinant nucleic acid molecule.
7. (Currently Amended) The composition of ~~any one of claims 1-6~~ claim 1, wherein the nucleic acid molecule is a vector, particularly an expression vector.
8. (Currently Amended) The composition of ~~any one of claims 1-5~~ claim 1, wherein the polypeptide is a recombinant polypeptide.
9. (Original) The composition of claim 8, wherein said recombinant polypeptide is a fusion polypeptide.
10. (Currently Amended) The composition of ~~any one of claims 1-7~~ claim 1, wherein said nucleic acid molecule is selected from hybridization probes, primers or/and anti-sense oligonucleotides.
11. (Currently Amended) The composition of ~~any one of claims 1-10~~ claim 1 which is a diagnostic composition.

12. (Currently Amended) The composition of ~~any one of claims 1-10~~ claim 1 which is a therapeutic composition.
13. (Currently Amended) The composition of ~~any one of claims 1-12~~ claim 1 for the manufacture of an agent for detecting or/and verifying, for the treatment, alleviation or/and prevention of metabolic diseases or dysfunctions, including metabolic syndrome, obesity, or/and diabetes, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, gallstones, or liver fibrosis in cells, cell masses, organs or/and subjects.
14. (Currently Amended) The composition of ~~any one of claims 1-13~~ claim 1 for application in vivo.
15. (Currently Amended) The composition of ~~any one of claims 1-13~~ claim 1 for application in vitro.
16. (Original) Use of a nucleic acid molecule encoding a *fwd*, *Pp2C1*, *Adk3*, *CG3860*, *Cdk4*, *CG7134*, or *Eip75B* homologous protein or/and an isoform, a functional fragment or/and a variant thereof, in particular a nucleic acid molecule as described in Table 1, particularly of a nucleic acid molecule according to claim 3 (a), (b), or (c), or/and a polypeptide encoded thereby or/and a functional fragment or/and a variant of said nucleic acid molecule or

said polypeptide or/and a modulator/effector of said nucleic acid molecule or said polypeptide for the manufacture of a medicament for treatment of obesity, diabetes, or/and metabolic syndrome for controlling the function of a gene or/and a gene product which is influenced or/and modified by a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide.

17. (Original) Use of the nucleic acid molecule encoding a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous protein or/and an isoform, a functional fragment or/and a variant thereof, in particular a nucleic acid molecule as described in Table 1, particularly of a nucleic acid molecule according to claim 3 (a), (b), or (c), or/and a polypeptide encoded thereby or/and a functional fragment or/and a variant of said nucleic acid molecule or said polypeptide, or/and a modulator/effector of said nucleic acid molecule or said polypeptide for identifying substances capable of interacting with a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide, particularly with a polypeptide according to claim 3.
18. (Original) A non-human transgenic animal exhibiting a modified expression of a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide, particularly of a polypeptide according to claim 3.
19. (Currently Amended) The animal of claim 18, wherein the expression of the *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous

polypeptide, ~~particularly of a polypeptide according to claim 3~~, is increased or/and reduced.

20. (Original) A recombinant host cell exhibiting a modified expression of a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide, particularly of a polypeptide according to claim 3.
21. (Original) The cell of claim 20 which is a human cell.
22. (Original) A method of identifying a (poly)peptide involved in the regulation of energy homeostasis or/and metabolism of triglycerides in a mammal comprising the steps of
  - (a) contacting a collection of (poly)peptides with a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide, particularly with a polypeptide according to claim 3, or a functional fragment thereof under conditions that allow binding of said (poly)peptides;
  - (b) removing (poly)peptides which do not bind and
  - (c) identifying (poly)peptides that bind to said *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide.
23. (Original) A method of screening for an agent which modulates/effects the interaction of a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B*

homologous polypeptide, particularly of a polypeptide according to claim 3, with a binding target/agent, comprising the steps of

- (a) incubating a mixture comprising
  - (aa) a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide, particularly a polypeptide according to claim 3, or a functional fragment thereof;
  - (ab) a binding target/agent of said polypeptide or functional fragment thereof; and
  - (ac) a candidate agentunder conditions whereby said polypeptide or functional fragment thereof specifically binds to said binding target/agent at a reference affinity;
- (b) detecting the binding affinity of said polypeptide or functional fragment thereof to said binding target to determine an affinity for the agent; and
- (c) determining a difference between affinity for the agent and reference affinity.

24. (Original) A method of screening for an agent, which modulates/effects the activity of a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide, comprising the steps of

- (a) incubating a mixture comprising
  - (aa) a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous polypeptide or a functional fragment thereof; and

- (ab) a candidate agent  
under conditions whereby said polypeptide or functional fragment thereof has a reference activity,
- (b) detecting the activity of said polypeptide or functional fragment thereof to determine an activity in presence of the agent; and
- (c) determining a difference between the activity in presence of the agent and the reference activity.

25. (Currently Amended) A method of producing a composition comprising mixing the (poly)peptide identified by the method of claim 22 ~~or the agent identified by the method of claim 23 or 24~~ with a pharmaceutically acceptable carrier, diluent, or/and additive.

26. (Original) The method of claim 25 wherein said composition is a pharmaceutical composition for preventing, alleviating, or/and treating of metabolic diseases or dysfunctions, including metabolic syndrome, obesity, or/and diabetes, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, gallstones, or liver fibrosis.

27. (Currently Amended) Use of a (poly)peptide as identified by the method of claim 22 ~~or of an agent as identified by the method of claim 23 or 24~~ for the preparation of a pharmaceutical composition for the treatment, alleviation



or/and prevention of metabolic diseases or dysfunctions, including metabolic syndrome, obesity, or/and diabetes, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, gallstones, or liver fibrosis.

28. (Currently Amended) Use of a nucleic acid molecule as defined in ~~any of claims 1-6 or 10~~ claim 1 for the preparation of a medicament for the treatment, alleviation or/and prevention of metabolic diseases or dysfunctions, including obesity, diabetes, or/and metabolic syndrome, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, osteoarthritis, gallstones, or liver fibrosis.
29. (Currently Amended) Use of a polypeptide as defined in ~~any one of claims 1 to 6, 8 or 9~~ claim 1 for the preparation of a medicament for the treatment, alleviation or/and prevention of metabolic diseases or dysfunctions, including obesity, diabetes, or/and metabolic syndrome, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, osteoarthritis, gallstones, or liver fibrosis.
30. (Original) Use of a vector as defined in claim 7 for the preparation of a medicament for the treatment, alleviation or/and prevention of metabolic

diseases or dysfunctions, including obesity, diabetes, or/and metabolic syndrome, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, osteoarthritis, gallstones, or liver fibrosis.

31. (Currently Amended) Use of a host cell as defined in claim 20 ~~or 21~~ for the preparation of a medicament for the treatment, alleviation or/and prevention of metabolic diseases or dysfunctions, including obesity, diabetes, or/and metabolic syndrome, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, osteoarthritis, gallstones, or liver fibrosis.
32. (Original) Use of a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous nucleic acid molecule or/and of a fragment thereof for the production of a non-human transgenic animal which over- or under-expresses the *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous gene product.
33. (Original) Kit comprising at least one of
- (a) a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous nucleic acid molecule or/and a fragment thereof;

- (b) a *fwd*, *Pp2C1*, *Adk3*, CG3860, *Cdk4*, CG7134, or *Eip75B* homologous amino acid molecule or/and a functional fragment or/and an isoform thereof;
- (c) a vector comprising the nucleic acid of (a);
- (d) a host cell comprising the nucleic acid of (a) or the vector of (c);
- (e) a polypeptide encoded by the nucleic acid of (a), expressed by the vector of (c) or the host cell of (d);
- (f) a fusion polypeptide encoded by the nucleic acid of (a);
- (g) an antibody, an aptamer or another modulator/effector of the nucleic acid of (a) or the polypeptide of (b) , (e) , or (f) or/and
- (h) an anti-sense oligonucleotide of the nucleic acid of (a).

34. (New) A method of producing a composition comprising mixing the agent identified by the method of claim 23 with a pharmaceutically acceptable carrier, diluent, or/and additive.

35. (New) Use of an agent as identified by the method of claim 23 for the preparation of a pharmaceutical composition for the treatment, alleviation or/and prevention of metabolic diseases or dysfunctions, including metabolic syndrome, obesity, or/and diabetes, as well as related disorders such as eating disorder, cachexia, pancreatitis, hypertension, coronary heart disease, hypercholesterolemia, dyslipidemia, gallstones, or liver fibrosis.